## Amendments to the Claims

1. (Currently amended) A method of treating an infection caused by herpesvirinae in a patient in need thereof comprising administering to said patient an effective amount of at least one compound according to Compounds comprising the chemical structure

$$X_3$$
Et  $Y_1$ 

wherein  $X_1$ ,  $X_2$ , and  $X_3$  are selected from the group consisting of O, N, and S; wherein  $Y_1$  and  $Y_2$  are selected from the group consisting of O, N, and S; and wherein Z is selected from the group consisting of F, Cl, and Br.

- 2. (Currently amended) A method as defined in claim 1, wherein the patient is administered an effective amount of at least one P pharmaceutically acceptable salts of the compounds of Claim 1.
- 3. (Currently amended) A method as defined in claim 1, wherein said Ccompounds having have substantially identical spatial occupation, physiochemical and electrochemical properties as the compounds of Claim 1.
- 4. (Currently amended) A method of treating an infection caused by herpesvirinae in a patient in need thereof comprising administering to said patient an effective amount of a compound consisting essentially of the chemical structure

or a pharmaceutically acceptable salt thereof.

- 5. (Currently amended) A pharmaceutical composition method as defined in claim 1 consisting essentially of comprising the administration of an effective amount of a compound of claim 1 and a pharmaceutically acceptable carrier.
- 6. (Currently amended) A method of treating an infection caused by herpesvirinae in a patient in need thereof comprising administering to said patient an effective amount of at least one compound having the three-dimensional structure characterized by the atomic structure coordinates of Table 5, said compound having less than a 10% difference in the internal coordinates after minimalization with the MM2 force field.
- 7. (New) A method according to claim 1, wherein the infection is caused by HSV-1 or HSV-2.